

**Figure S10: CD117- and GluA4-targeted NiV-LV.** (**A**) Scatter dot blot of fluorescence intensities of Fig 9A. Mean fluorescence intensities of surface expression of CD117 and CD117short on HT1080 are shown. Surface expression was analyzed using a CD117-

specific antibody. Parental HT1080 cells served as negative control (mock) (n=4; mean ± standard deviations (SD) are shown; \*\*, P<0.01 by unpaired t-test). (B) Scatter dot blot of fluorescence intensities of Fig 9C. Mean fluorescence intensities of surface binding of Fc-SCF to HT1080-CD117 and HT1080-CD117short are shown. Surface binding was analyzed using a FITC coupled anti-Fc antibody. Parental HT1080 cells served as negative control (mock) (n=3; mean ± standard deviations (SD) are shown; ns, not significant by unpaired t-test). (C) Exemplary flow cytometry blot of surface expression of NiV-Gc∆34<sup>CD117</sup>mut4 (black line). HEK-293T cells were transiently transfected with plasmids encoding the glycoprotein and compared to mock transfected cells (filled curve) as determined by flow cytometry. Cells were stained with PE coupled anti-His antibody. One representative out of three experiments is shown. (D) Scatter dot blot of fluorescence intensities in (C). Mean fluorescence intensities of surface expression of the CD117-targeted G variant on HEK-293T cells transiently transfected with the corresponding expression plasmid compared to mock transfected cells as determined by flow cytometry. Cells were stained with PE-coupled anti-His antibody (n=3; mean ± standard deviations (SD) are shown). (E) Scatter dot blot of fluorescence intensities of Fig 9B. Mean fluorescence intensities of surface expression of GluA4 and GluA4short on HT1080 are shown. Surface expression was analyzed using a myc-tag specific antibody. Parental HT1080 cells served as negative control (mock) (n=3; mean ± standard deviations (SD) are shown; \*\*, P<0.01 by unpaired t-test). (**F**) Exemplary flow cytometry blot of surface expression of NiV-Gc∆34 GluA4 mut4 (black line). HEK-293T cells were transiently transfected with plasmids encoding the glycoprotein and compared to mock transfected cells (filled curve) as determined by flow cytometry. Cells were stained with PE coupled anti-His antibody. One representative out of three experiments is shown. (G) Scatter dot blot of fluorescence intensities in (**F**). Mean fluorescence intensities of surface expression of GluA4-targeted G variant on HEK-293T cells transiently transfected with the corresponding expression plasmid compared to mock transfected cells as determined by flow cytometry. Cells were stained with PE-coupled anti-His antibody (n=3; mean ± standard deviations (SD) are shown).